Regulation (EU) 2017/746 of the European Parliament and of the Council of 5 April 2017 on in vitro diagnostic medical devices and repealing Directive 98/79/EC and Commission Decision 2010/227/EU (Text with EEA relevance) (revoked)

CHAPTER VII

POST-MARKET SURVEILLANCE, VIGILANCE AND MARKET SURVEILLANCE

Section 3

Market surveillance

Article 89

Evaluation of devices suspected of presenting an unacceptable risk or other non-compliance

. . . . . .

Textual Amendments applied to the whole legislation

F1 Regulation revoked (31.12.2020) by The Medical Devices Regulations 2002 (S.I. 2002/618), **reg. 4P** (as inserted by S.I. 2019/791, regs. 1(1), 3(7) (as amended by S.I. 2020/1478, regs. 1(3), Sch. 2 paras. 2, 9(m)); 2020 c. 1, Sch. 5 para. 1(1))

Status:
This version of this provision no longer has effect.

Changes to legislation:
There are currently no known outstanding effects for the Regulation (EU) 2017/746 of the European Parliament and of the Council, Article 89.